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NEWS	3	JAN		CAS patent coverage enhanced to include exemplified
		01111		prophetic substances
NEWS	4	JAN	28	USPATFULL, USPAT2, and USPATOLD enhanced with new custom IPC display formats
NEWS	5	JAN	28	MARPAT searching enhanced
NEWS	6	JAN	28	USGENE now provides USPTO sequence data within 3 days
				of publication
NEWS	7	JAN		TOXCENTER enhanced with reloaded MEDLINE segment
NEWS	8	JAN		MEDLINE and LMEDLINE reloaded with enhancements
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NEWS	13	FEB	29	WPINDEX/WPIDS/WPIX enhanced with ECLA and current
				U.S. National Patent Classification
NEWS	14	MAR	31	IFICDB, IFIPAT, and IFIUDB enhanced with new custom
				IPC display formats
NEWS	15	MAR	31	CAS REGISTRY enhanced with additional experimental
				spectra
NEWS	16	MAR	31	CA/CAplus and CASREACT patent number format for U.S.
				applications updated
NEWS	17	MAR	31	LPCI now available as a replacement to LDPCI
NEWS	18	MAR	31	EMBASE, EMBAL, and LEMBASE reloaded with enhancements
NEWS	19	APR	04	STN AnaVist, Version 1, to be discontinued
NEWS	20	APR	15	WPIDS, WPINDEX, and WPIX enhanced with new
				predefined hit display formats
NEWS	21	APR	28	EMBASE Controlled Term thesaurus enhanced
NEWS	22	APR	28	IMSRESEARCH reloaded with enhancements
NEWS	23	MAY	30	INPAFAMDB now available on STN for patent family
				searching
NEWS	24	MAY	30	DGENE, PCTGEN, and USGENE enhanced with new homology
				sequence search option
NEWS	25	JUN	0.6	EPFULL enhanced with 260,000 English abstracts
NEWS		JUN		KOREAPAT updated with 41,000 documents
NEWS		JUN		USPATFULL and USPAT2 updated with 11-character
111110		0011		patent numbers for U.S. applications
NEWS	28	JUN	19	CAS REGISTRY includes selected substances from
112110	20	0011		web-based collections
NEWS	29	JUN	25	CA/CAplus and USPAT databases updated with IPC
MEMO	23	0.014	-5	reclassification data
NEWS	3.0	JUN	3.0	AEROSPACE enhanced with more than 1 million U.S.
MEMO	50	OON	50	patent records
NEWS	21	JUN	20	EMBASE, EMBAL, and LEMBASE updated with additional
NEWS	21	OON	50	EMDAGE, EMDAE, AND EEMDAGE UPUGLED WILL ADDITIONAL

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=> file medline, biosis, wpids, uspatful, hcaplus

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=> s (inflammation and ferritin)

L1 3209 (INFLAMMATION AND FERRITIN)

=> s l1 and biliverdin

L2 72 L1 AND BILIVERDIN

=> s 12 and dosage

L3 31 L2 AND DOSAGE

=> s 13 and (dextran)

.4 19 L3 AND (DEXTRAN)

=> s 13 and (sesferoxamine

UNMATCHED LEFT PARENTHESIS 'AND (SESFEROXAM'

The number of right parentheses in a query must be equal to the number of left parentheses.

=> s 13 and (desferoxamine) L5 12 L3 AND (DESFEROXAMINE) => s 15 and 14 L6 12 L5 AND L4

=> d 16 ti abs ibib tot

L6 ANSWER 1 OF 12 WPIDS COPYRIGHT 2008 THOMSON REUTERS on STN
T Treating inflammation involves determining the level of heme
oxygenase-1 activity in response to a stimulus; and administering
anti-inflammatory agent, and composition having carbon monoxide,
bilirubin, heme oxygenase-1, and/or apoferritin
AN 2008-C63174 [19] WPIDS

AB WO 2008008513 A2 UPAB: 20080318

NOVELTY - Treating inflammation involves determining the patient's level of heme oxygenase-1 (HO-1) activity, expression, or induction in response to a stimulus, or determining an allele of a polymorphism in HO-1 promoter; and administering first pharmaceutical composition comprising an anti-inflammatory agent; and a second pharmaceutical composition comprising carbon monoxide, HO-1, bilirubin, iron dextran, apoferritin (if HO-1 activity, expression or induction in response to stimulus is determined to be reduced compared to reference standard, or if the HO-1 promoter comprises specified allele).

DETAILED DESCRIPTION - Treating (M1) inflammation

involves determining the patient's level of heme oxygenase-1 (HO-1) activity, expression, or induction in response to a stimulus, or determining an allele of a polymorphism in HO-1 promoter; and administering first pharmaceutical composition comprising an anti-inflammatory agent; and a second pharmaceutical composition comprising carbon monoxide (CO), a CO-releasing compound, HO-1, bilirubin, biliverdin, ferritin, iron, desferoxamine, salicylaldehyde isonicotinoyl hydrazone, iron dextran, and/or apoferritin (if HO-1 activity, expression or induction in response to a stimulus is determined to be reduced compared to a reference standard, or if the HO-1 promoter comprises a specified allele). An INDEPENDENT CLAIM is included for potentiating (M2) the response of a patient to a pharmaceutical agent, involving identifying a first pharmaceutical agent that is potentiated by a second treatment which induces HO-1 or apoferritin or increases the level of expression of HO-1 or apoferritin in the patient by administering a second pharmaceutical composition comprising HO-1, CO, CO-releasing compound, bilirubin, biliverdin , ferritin, iron, desferoxamine, salicylaldehyde isonicotinoyl hydrazone, iron dextran, and/or apoferritin; administering the first pharmaceutical agent; and allowing the second treatment. PHARMACEUTICALS - Preferred Components: The anti-inflammatory

pharmaceutical agent is immunosuppresant.

ACTIVITY - Antiinflammatory; Antiasthmatic; Respiratory-Gen.;
Hypotensive; Cardiovascular-Gen.; Vasotropic; Cerebroprotective;
Antiarteriosclerotic; Cardiant; Nephrotropic; Uropathic; Hepatotropic;
Virucide, Antianqioqenic; Gastrointestinal-Gen.; Antiarthritic;
Antirheumatic; Neuroprotective; Dermatological; Immunosuppressive;
Cytostatic; Vulnerary; Nootropic; Antiparkinsonian; Hemostatic;
Antibacterial; Analgesic; Gynecological; Endocrine-Gen.; Anti-HIV;
Antiallerqic. The efficacy of heme oxygenase-l was tested for anti-inflammatory effect. Mouse macrophage cell lines were stably

agent is selected from statins, adenosine, cyclooxygenase inhibitors, probucol, steroids, or prostaglandins. In method (M2) the first

transfected with either heme oxygenase-1 (HO-I) or empty plasmid (NEO). The resulting cell lines were treated with adenosine (100 muM) or 5'-(N-ethylcarboxamido) adenosine (NECA) (10 muM) 30 minutes prior to stimulation with lipopolysaccharide (LPS) (1 ng/ml). After 4 hours, the supernatant of each group was analyzed for tumor necrosis factor (TNF)-I. Overexpression of HO-1 gave approximately1.5 to 2-fold greater inhibition of TNF-I secretion compared to the vector control. Thus overexpression of HO-1 augmented the effect of the anti-inflammatory agents adenosine and NECA on TNF-I, produced by lipopolysaccharide (LPS) activated macrophages. MECHANISM OF ACTION - Heme oxygenase 1 stimulator; Apoferritin

stimulator. No biological data given. USE - For treating inflammation (particularly associated with asthma, adult respiratory distress syndrome, interstitial pulmonary fibrosis, pulmonary emboli, chronic obstructive pulmonary disease, primary pulmonary hypertension, chronic pulmonary emphysema, congestive heart failure, peripheral vascular disease, stroke, atherosclerosis, ischemia-reperfusion injury, heart attack, glomerulonephritis, conditions involving inflammation of the kidney, infection of the genitourinary tract, viral hepatitis, toxic hepatitis, cirrhosis, ileus, necrotizing enterocolitis, inflammatory bowel disease, rheumatoid arthritis, multiple sclerosis, systemic lupus erythematosus, cancer, wounds, Alzheimer's disease, Parkinson's disease, graft versus host disease, and hemorrhagic, septic, or anaphylactic shock); inflammation of the heart, respiratory tract, liver, spleen, brain, joint, skin, castrointestinal tract and/or kidney (claimed), pain, reproductive disorders, e.g. impotence, premature uterine contractions, premature deliveries and menstrual cramps, amoebic dysentery, pneumonia (bacterial or viral), inflammatory states due to immunodeficiency e.g. due to infection with HIV; hypersensitivities; and for treating unwanted angiogenesis.

ADVANTAGE - The second composition induces HO-1 or increasing the level of expression of HO-1; induces apoferritin or increases the level of expression of apoferritin in the patient, and effectively treat inflammation. WPIDS

ACCESSION NUMBER: 2008-C63174 [19]

DOC. NO. CPI: C2008-079637 [19]

TITLE: Treating inflammation involves determining the

level of heme oxygenase-1 activity in response to a stimulus; and administering anti-inflammatory agent, and

composition having carbon monoxide, bilirubin, heme oxygenase-1, and/or apoferritin

DERWENT CLASS: B05

INVENTOR . BACH F H; HASCHEMI A; OTTERBEIN L E

PATENT ASSIGNEE: (BETH-N) BETH ISRAEL DEACONESS MEDICAL CENT

COUNTRY COUNT: 120

PATENT INFO ABBR.:

PA	TENT	NO	KIND	DATE	WEEK	LA	PG	MAIN	IPC
		8008513 8008513		20080117 20080306	(200819)* (200819)	EN EN	49[4]		

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 2008008513	A2	WO 2007-US160	32 20070712

PRIORITY APPLN. INFO: US 2006-830480P 20060713

L6 ANSWER 2 OF 12 WPIDS COPYRIGHT 2008 THOMSON REUTERS on STN

II Use of heme oxygenase-1 and heme degradation products for e.g. reducing inflammation, organ transplantation and treating e.g. cellular proliferative disorder

AN 2003-903222 [82] WPIDS

AB WO 2003088748 A1 UPAB: 20060121

NOVELTY - Reducing (M1) inflammation involves:

- administration of at least one treatment selected from inducing ferritin;
 - (2) expressing ferritin; and
- (3) administering a pharmaceutical composition (C1) comprising heme oxygenase-1 (HO-1), bilirubin, biliverdin, ferritin,

iron, desferoxamine, salicylaldehyde isonicotinoyl hydrazone, iron dextran, or apoferritin.

DETAILED DESCRIPTION - INDEPENDENT CLAIMS are also included for:

(1) A method (M2) of transplanting an organ by three different ways (M2a, M2b, M2c). (M2a) involves (ia) administration of at least one of the treatments to a donor to enhance survival or function of the organ after the transplantation, (ib) obtaining an organ from the donor, and (ic) transplanting the organ into a recipient. (M2b) involves (iia) administering to an organ of a donor ex vivo at least one of the treatments, and (iib) transplanting the organ into a recipient. (M2c) involves (iiia) transplanting the organ into a recipient, and (iib) administering at least one of the treatments to the recipient; and

(2) A method (M3) of performing angioplasty and vascular surgery involving performing angioplasty and vascular surgery, respectively, and administering at least one of the treatments.

ACTIVITY - Anti-inflammatory; Antiulcer; Antiasthmatic; Tranquilizer; Respiratory-Gen.; Thrombolytic; Hypotensive; Cardiovascular Gen.; Cerebroprotective; Antiarteriosclerotic; Vasotropic; Cardiant; Nephrotropic; Hepatotropic; Virucide; Gastrointestinal Gen.; Antirheumatic; Antiarthritic; Vulnerary; Neuroprotective; Nootropic; Antiparkinsonian; Immunosuppressive; Antibacterial; Uropathic; Cytostatic; Gynecological.

The anti-inflammatory efficacy of biliverdin was evaluated in an animal model of endotoxic shock. Administration of endotoxin in male Sprague-Dawley rats resulted in lung inflammation, neutrophil accumulation, and increased levels of tumor necrosis factor-alpha (TNF-alpha) in the serum. The rats were administered with biliverdin (50 micromol/kg) 17 hours prior to, and 8 hours after endotoxin administration. Serum level of TNF-alpha was measured by ELISA kits, and total cell counts was determined by differential analysis. The results showed that biliverdin reduced levels of TNF-alpha; levels of neutrophils and protein accumulation in the airspace; and also increased the levels of anti-inflammatory cytokine IL-10.

MECHANISM OF ACTION - None given.

USE - The treatment is useful for reducing and treating inflammation of the heart, lung, liver, spleen, brain skin and kidney; inflammatory condition (e.g. amoebic dysentery, bacillary dysentery, schistosomiasis, campylobacter enterocolitis, yersinia enterocolitis, enterobbis vermicularis, radiation enterocolitis, ischaemic colitis, eosinophilic gastroenteritis, ulcerative colitis, indeterminate colitis, and Crohn's disease) localized in the gastrointestinal tract); asthma; adult respiratory distress syndrome; interstitial pulmonary fibrosis; pulmonary emboli; chronic obstructive pulmonary disease; primary pulmonary hypertension; chronic pulmonary emphysema; congestive heart failure; peripheral vascular disease; stroke; atherosclerosis; ischemia-reperfusion injury; heart attacks; glomevulonephritis; nephrotic disorders; infection of the genitourinary tract; viral and toxic hepatitis; cirrhosis; ileus; necrotizing enterocolitis; specific and

non-specific inflammatory bowel disease; rheumatoid arthritis, deficient wound healing, Alzheimer's disease, Parkinson's disease, graft versus host disease, and hemorrhagic, septic, or anaphylactic shock; cellular proliferative and differentiative disorder. For reducing the effects of ischemia. For transplanting organs and performing angioplasty or vascular surgery (all claimed). Also for treating other autoimmune diseases and reproductive disorders.

ADVANTAGE - The heme-oxygenase-1 and the heme degradation products attenuate inflammation and suppress the damage associated with

ischemia.

ACCESSION NUMBER: 2003-903222 [82] WPIDS DOC. NO. CPI: C2003-256695 [82]

DOC. NO. NON-CPI: N2003-721263 [82]

TITLE: Use of heme oxygenase-1 and heme degradation products for e.g. reducing inflammation, organ

transplantation and treating e.g. cellular proliferative

disorder

DERWENT CLASS: B04; B05; S03

INVENTOR: BACH F H; BERBERAT P O; ROBSON S C

PATENT ASSIGNEE: (BETH-N) BETH ISRAEL DEACONESS MEDICAL CENT; (BACH-I)
BACH F H; (BERB-I) BERBERAT P O; (ROBS-I) ROBSON S C

COUNTRY COUNT: 102

PATENT INFO ABBR.:

PAT	ENT NO	KIN	DATE	WEEK	LA	PG	MAIN	IPC
	2003088748			(200382)*		56 [27]		
	2003226366 1499186		20031103		EN EN			
	2005522521		20050728		JA	59		
US	20060003922	A1	20060105	(200603)	EN			

APPLICATION DETAILS:

PATENT NO KIND	APPLICATION DATE
WO 2003088748 A1	WO 2003-US11411 20030415
AU 2003226366 A1	AU 2003-226366 20030415
EP 1499186 A1	EP 2003-746978 20030415
JP 2005522521 W	JP 2003-585506 20030415
EP 1499186 A1	WO 2003-US11411 20030415
JP 2005522521 W	WO 2003-US11411 20030415
US 20060003922 A1 Provis	ional US 2002-372762P 20020415
US 20060003922 A1	WO 2003-US11411 20030415
US 20060003922 A1	US 2005-511612 20050805

FILING DETAILS:

PA'	TENT NO	KIND	PA	TENT NO	
EP	2003226366 A1 1499186 A1 2005522521 W	Based Based Based	on WO	2003088748 2003088748 2003088748	A
PRIORITY	APPLN. INFO:	US 2002-372		20415	

L6 ANSWER 3 OF 12 USPATFULL on STN

TI Carbon monoxide improves outcomes in tissue and organ transplants and suppresses apoptosis

US 2005-511612 20050805

AB The present invention features methods for transplanting organs, tissues and individual cells. Also featured are methods for maintaining cells in vitro and for enhancing survival and/or function of cells following transplantation. The methods include the administration of carbon monoxide in an amount sufficient to enhance cell survival and/or function.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

2007:230812 USPATFULL ACCESSION NUMBER:

TITLE: Carbon monoxide improves outcomes in tissue and organ

transplants and suppresses apoptosis

INVENTOR(S): Bach, Fritz H., Manchester-by-the Sea, MA, UNITED

STATES

Otterbein, Leo E., New Kensington, PA, UNITED STATES

Soares, Miguel P., Boston, MA, UNITED STATES

Tobiasch, Edda M., Bonn, GERMANY, FEDERAL REPUBLIC OF Gose, Jeanne, Manchester-by-the Sea, MA, UNITED STATES

PATENT ASSIGNEE(S): Beth Israel Deaconess Medical Center, Inc., a

Massachusetts corporation (U.S. corporation)

NUMBER KIND DATE PATENT INFORMATION: US 20070202083 A1 20070830 US 2006-401722 A1 20060410 (11)

APPLICATION INFO.: RELATED APPLN. INFO.: Division of Ser. No. US 2002-177930, filed on 21 Jun

2002, GRANTED, Pat. No. US 7238469

DATE NUMBER -----PRIORITY INFORMATION:

US 2001-300289P 20010621 (60) US 2001-334340P 20011129 (60) US 2001-337974P 20011207 (60)

DOCUMENT TYPE: Utility FILE SEGMENT:

APPLICATION

LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, P.O. BOX 1022, MINNEAPOLIS, MN, 55440-1022, US

NUMBER OF CLAIMS: 29

NUMBER OF CLAIMS: 25 EXEMPLARY CLAIM: 1-47 NUMBER OF DRAWINGS: 31 Drawing Page(s)

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 4 OF 12 USPATFULL on STN

TI Spinner preparation machine and cavity resonator

AB The present invention relates to the treatment of disorders using heme oxygenase-1 and heme degradation products.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2006:4464 USPATFULL

TITLE: Spinner preparation machine and cavity resonator INVENTOR(S): Bach, Fritz H., Manchester-by-the-sea, MA, UNITED

Berberat, Pascal O., Heidelberg, GERMANY, FEDERAL

REPUBLIC OF

Robson, Simon C., Weston, MA, UNITED STATES

NUMBER KIND DATE PATENT INFORMATION: US 20060003922 A1 20060105 APPLICATION INFO.: US 2003-511612 A1 20030415 (10) WO 2003-US11411 20030415 NUMBER DATE

PRIORITY INFORMATION: US 2002-372762P 20020415 (60)

DOCUMENT TYPE: Utility

FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, P.O. BOX 1022, MINNEAPOLIS, MN,

55440-1022, US NUMBER OF CLAIMS: 58

EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 27 Drawing Page(s) LINE COUNT: 3083

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

1.6 ANSWER 5 OF 12 USPATFULL on STN

ΤI Methods of treating angiogenesis, tumor growth, and metastasis

The present invention relates to a method of treating cancer or unwanted AB angiogenesis in a patient, which includes administering a pharmaceutical composition that includes carbon monoxide to the patient.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

2004:326952 USPATFULL ACCESSION NUMBER:

TITLE: Methods of treating angiogenesis, tumor growth, and

metastasis

Otterbein, Leo E., New Kensington, PA, UNITED STATES INVENTOR(S): Choi, Augustine M. K., Pittsburgh, PA, UNITED STATES

NUMBER KIND DATE PATENT INFORMATION: US 20040258772 A1 20041223 US 2003-455564 A1 20030605 (10) APPLICATION INFO.:

NUMBER DATE

PRIORITY INFORMATION: US 2002-386561P 20020605 (60)

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICAT APPLICATION

LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, 225 FRANKLIN ST, BOSTON, MA,

02110 NUMBER OF CLAIMS: 71

EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 10 Drawing Page(s)
LINE COUNT: 1303

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 6 OF 12 USPATFULL on STN 1.6

TI Treatment for hemorrhagic shock

AB The present invention relates to methods and compositions of treating patients suffering from, or at risk for, hemorrhagic shock. The treatment includes administering to the patient a pharmaceutical composition that includes carbon monoxide.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

2004:291849 USPATFULL ACCESSION NUMBER: TITLE:

Treatment for hemorrhagic shock

INVENTOR(S): Billiar, Timothy R., Nevillewood, PA, UNITED STATES Choi, Augustine M.K., Pittsburgh, PA, UNITED STATES McCloskey, Carol A., Pittsburgh, PA, UNITED STATES Otterbein, Leo E., New Kensington, PA, UNITED STATES Zuckerbraun, Brian, Pittsburgh, PA, UNITED STATES

NUMBER KIND DATE PATENT INFORMATION: US 20040228930 A1 20041118

US 2003-676280 APPLICATION INFO.: A1 20030930 (10) NUMBER DATE

PRIORITY INFORMATION: US 2002-424804P 20021107 (60)

DOCUMENT TYPE: Utility

FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, 225 FRANKLIN ST, BOSTON, MA,

02110 NUMBER OF CLAIMS: 5.4

EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 9 Drawing Page(s)

TIME COUNT: 1154 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 7 OF 12 USPATFULL on STN

ΤI Pharmaceutical use of nitric oxide, heme oxygenase-1 and products of heme degradation

The present invention relates to the treatment of disorders using nitric AB oxide (NO), heme oxygenase-1 (HO-1) and heme degradation products such as carbon monoxide (CO), biliverdin, bilirubin and iron.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2004:171542 USPATFULL

TITLE: Pharmaceutical use of nitric oxide, heme oxygenase-1

and products of heme degradation

INVENTOR(S): Bach, Fritz H., Manchester-by-the-sea, MA, UNITED

Otterbein, Leo E., New Kensington, PA, UNITED STATES

NUMBER KIND DATE ______ PATENT INFORMATION: US 20040131703 A1 20040708 APPLICATION INFO.: US 2003-600182 A1 20030620 (10)

NUMBER DATE

PRIORITY INFORMATION: US 2002-390457P 20020621 (60)

DOCUMENT TYPE: FILE SEGMENT: Utility

APPLICATION

LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, 225 FRANKLIN ST, BOSTON, MA,

02110 NUMBER OF CLAIMS: 23

EXEMPLARY CLAIM:

1 8 Drawing Page(s) NUMBER OF DRAWINGS:

LINE COUNT:

2300

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 8 OF 12 USPATFULL on STN

TI Methods of treating hepatitis

AB The present invention relates to a method of treating hepatitis in a patient, which includes administering a pharmaceutical composition that includes carbon monoxide to the patient.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2004:69639 USPATFULL TITLE . Methods of treating hepatitis INVENTOR(S): Otterbein, Leo E., New Kensington, PA, UNITED STATES Choi, Augustine M. K., Pittsburgh, PA, UNITED STATES

Zuckerbraun, Brian, Pittsburgh, PA, UNITED STATES

NUMBER KIND DATE US 20040052866 A1 20040318 PATENT INFORMATION: APPLICATION INFO.: US 2003-439632 A1 20030516 (10)

NUMBER DATE

PRIORITY INFORMATION: US 2002-381527P 20020517 (60)

DOCUMENT TYPE: Utility

FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, 225 FRANKLIN ST, BOSTON, MA, 02110

NUMBER OF CLAIMS: 26

EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 21 Drawing Page(s)

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L6 ANSWER 9 OF 12 USPATFULL on STN

TΙ Methods of treating necrotizing enterocolitis

AB The present invention relates to a method of treating necrotizing enterocolitis in a patient, which includes administering a

pharmaceutical composition that includes carbon monoxide to the patient.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2004:7134 USPATFULL

TITLE: Methods of treating necrotizing enterocolitis

INVENTOR(S): Otterbein, Leo E., New Kensington, PA, UNITED STATES Zuckerbraun, Brian, Pittsburgh, PA, UNITED STATES

NUMBER KIND DATE ______ PATENT INFORMATION: US 20040005367 A1 20040108 APPLICATION INFO.: US 2003-413817 A1 20030415 (10)

NUMBER DATE

PRIORITY INFORMATION: US 2002-372599P 20020415 (60)

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, 225 FRANKLIN ST, BOSTON, MA,

02110

NUMBER OF CLAIMS: 34
EXEMPLARY CLAIM: 1
NUMBER OF DRAWINGS: 8 Drawing Page(s)
LINE COUNT: 1097

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 10 OF 12 USPATFULL on STN

Methods of treating ileus TI

AB The present invention relates to a method of treating ileus in a patient, which includes administering a pharmaceutical composition that includes carbon monoxide to the patient.

CAS INDEXING IS AVAILABLE FOR THIS PATENT. ACCESSION NUMBER: 2003:311902 USPATFULL TITLE . Methods of treating ileus INVENTOR(S): Otterbein, Leo E., New Kensington, PA, UNITED STATES Choi, Augustine M. K., Pittsburgh, PA, UNITED STATES

Moore, Beverley A., Pittsburgh, PA, UNITED STATES Bauer, Anthony J., Pittsburgh, PA, UNITED STATES

NUMBER KIND DATE ______ US 20030219497 A1 20031127 PATENT INFORMATION: APPLICATION INFO.: US 2003-371666 A1 20030221 (10)

NUMBER DATE

PRIORITY INFORMATION: US 2002-372652P 20020415 (60)

DOCUMENT TYPE: Utility

FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, 225 FRANKLIN ST, BOSTON, MA,

02110 NUMBER OF CLAIMS: 24

EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 16 D: 2256 16 Drawing Page(s) CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 11 OF 12 USPATFULL on STN

Methods of treating vascular disease

AB The present invention relates to a method of treating patients suffering from, or at risk for, intimal hyperplasia and/or arteriosclerosis. The

treatment includes administering a pharmaceutical composition that includes carbon monoxide to the patient.

CAS INDEXING IS AVAILABLE FOR THIS PATENT. ACCESSION NUMBER: 2003:311901 USPATFULL

Methods of treating vascular disease TITLE:

Otterbein, Leo E., New Kensington, PA, UNITED STATES INVENTOR(S): Choi, Augustine M. K., Pittsburgh, PA, UNITED STATES Bach, Fritz H., Mancester-by-the-sea, MA, UNITED STATES

Zuckerbraun, Brian, Pittsburgh, PA, UNITED STATES

NUMBER KIND DATE US 20030219496 A1 20031127 PATENT INFORMATION: US 7364757 B2 20080429
APPLICATION INFO.: US 2003-367277 A1 20030213 (10)

NUMBER DATE

PRIORITY INFORMATION: US 2002-356718P 20020213 (60)

DOCUMENT TYPE: Utility

FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, 225 FRANKLIN ST, BOSTON, MA,

02110

NUMBER OF CLAIMS: 49 EXEMPLARY CLAIM:

18 Drawing Page(s) 1841 NUMBER OF DRAWINGS:

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 12 OF 12 USPATFULL on STN

TI Carbon monoxide improves outcomes in tissue and organ transplants and suppresses apoptosis

AB The present invention features methods for transplanting organs, tissues and individual cells. Also featured are methods for maintaining cells in vitro and for enhancing survival and/or function of cells following transplantation. The methods include the administration of carbon monoxide in an amount sufficient to enhance cell survival and/or function.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2003:57074 USPATFULL

TITLE: Carbon monoxide improves outcomes in tissue and organ

transplants and suppresses apoptosis

INVENTOR(S): Bach, Fritz H., Manchester-by-the-Sea, MA, UNITED

STATES

Otterbein, Leo E., New Kensington, PA, UNITED STATES Soares, Miquel P., Boston, MA, UNITED STATES

Tobiasch, Edda M., Bonn, GERMANY, FEDERAL REPUBLIC OF Gose, Jeanne, Manchester-by-the-Sea, MA, UNITED STATES

		NUMBER	KIND	DATE	
PATENT INFORMATION:	US	20030039638	A1	20030227	
	US	7238469	B2	20070703	
APPLICATION INFO.:	US	2002-177930	A1	20020621	(10)

	NUMBER	DATE	
PRIORITY INFORMATION:	US 2001-300289P	20010621	(60)
	US 2001-334340P	20011129	(60)
	US 2001-337974P	20011207	(60)
DOCUMENT TYPE:	Utility		

FILE SEGMENT:

APPLICATION LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, 225 FRANKLIN ST, BOSTON, MA,

02110

NUMBER OF CLAIMS: 149

EXEMPLARY CLAIM: NUMBER OF DRAWINGS: 31 Drawing Page(s)

LINE COUNT: 3473

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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(FILE 'HOME' ENTERED AT 18:15:40 ON 06 JUL 2008)

FILE 'MEDLINE, BIOSIS, WPIDS, USPATFULL, HCAPLUS' ENTERED AT 18:15:59 ON 06 JUL 2008

3209 S (INFLAMMATION AND FERRITIN)

L2 72 S L1 AND BILIVERDIN L3 31 S L2 AND DOSAGE

19 S L3 AND (DEXTRAN) L4

L5 12 S L3 AND (DESFEROXAMINE)

L6 12 S L5 AND L4

=> s 16 and (colitis)

L7 4 L6 AND (COLITIS)

=> d 17 ti abs ibib tot

ANSWER 1 OF 4 WPIDS COPYRIGHT 2008 THOMSON REUTERS on STN

Use of heme oxygenase-1 and heme degradation products for e.g. reducing inflammation, organ transplantation and treating e.g. cellular proliferative disorder

AN

AB

NOVELTY - Reducing (M1) inflammation involves:

- administration of at least one treatment selected from inducing ferritin;
 - (2) expressing ferritin; and
- (3) administering a pharmaceutical composition (C1) comprising heme oxygenase-1 (HO-1), bilirubin, biliverdin, ferritin, iron, desferoxamine, salicylaldehyde isonicotinovl hydrazone,

iron dextran, or apoferritin.

DETAILED DESCRIPTION - INDEPENDENT CLAIMS are also included for:
(1) A method (M2) of transplanting an organ by three different ways

(M2a, M2b, M2c). (M2a) involves (ia) administration of at least one of the treatments to a donor to enhance survival or function of the organ after the treatments to a donor to enhance survival or function of the organ after the transplantation, (ib) obtaining an organ from the donor, and (ic) transplanting the organ into a recipient. (M2b) involves (iia) administering to an organ of a donor ex vivo at least one of the treatments, and (iib) transplanting the organ into a recipient. (M2c) involves (iiia) transplanting an organ from a donor into a recipient, and (iib) administering at least one of the treatments to the recipient; and

(2) A method (M3) of performing angioplasty and vascular surgery involving performing angioplasty and vascular surgery, respectively, and administering at least one of the treatments.

ACTIVITY - Anti-inflammatory; Antiulcer; Antiasthmatic; Tranquilizer; Respiratory-Gen.; Thrombolytic; Hypotensive; Cardiovascular Gen.; Cerebroprotective; Antiarteriosclerotic; Vasotropic; Cardiant; Nephrotropic; Hepatotropic; Virucide; Gastrointestinal Gen.; Antirheumatic; Antiatrhitic; Vulnerary; Neuroprotective; Nootropic; Antiparkinsonian; Immunosuppressive; Antibacterial; Uropathic; Cytostatic; Gynecological.

The anti-inflammatory efficacy of biliverdin was evaluated in an animal model of endotoxic shock. Administration of endotoxin in male Sprague-Dawley rats resulted in lung inflammation, neutrophil accumulation, and increased levels of tumor necrosis factor-alpha (TNF-alpha) in the serum. The rats were administered with biliverdin (50 micromol/kg) 17 hours prior to, and 8 hours after endotoxin administration. Serum Hevel of TNF-alpha was measured by ELISA kits, and total cell counts was determined by differential analysis. The results showed that biliverdin reduced levels of TNF-alpha; levels of neutrophils and protein accumulation in the airspace; and also increased the levels of anti-inflammatory cytokine IL-10.

MECHANISM OF ACTION - None given.

USE - The treatment is useful for reducing and treating inflammation of the heart, lung, liver, spleen, brain skin and kidney; inflammatory condition (e.g. amoebic dysentery, bacillary dysentery, schistosomiasis, campylobacter enterocolitis, versinia enterocolitis, enterobius vermicularis, radiation enterocolitis, ischaemic colitis, eosinophilic gastroenteritis, ulcerative colitis , indeterminate colitis, and Crohn's disease) localized in the qastrointestinal tract); asthma; adult respiratory distress syndrome; interstitial pulmonary fibrosis; pulmonary emboli; chronic obstructive pulmonary disease; primary pulmonary hypertension; chronic pulmonary emphysema; congestive heart failure; peripheral vascular disease; stroke; atherosclerosis; ischemia-reperfusion injury; heart attacks; glomerulonephritis; nephrotic disorders; infection of the genitourinary tract; viral and toxic hepatitis; cirrhosis; ileus; necrotizing enterocolitis; specific and non-specific inflammatory bowel disease; rheumatoid arthritis, deficient wound healing, Alzheimer's disease, Parkinson's disease, graft versus host disease, and hemorrhagic, septic, or anaphylactic shock; cellular proliferative and differentiative

disorder. For reducing the effects of ischemia. For transplanting organs and performing angioplasty or vascular surgery (all claimed). Also for treating other autoimmune diseases and reproductive disorders.

ADVANTAGE - The heme-oxygenase-1 and the heme degradation products attenuate inflammation and suppress the damage associated with ischemia.

ACCESSION NUMBER: 2003-903222 [82] WPIDS DOC. NO. CPI: C2003-256695 [82]

DOC. NO. NON-CPI: N2003-721263 [82]

TITLE: Use of heme oxygenase-1 and heme degradation products for

e.g. reducing inflammation, organ transplantation and treating e.g. cellular proliferative

disorder

DERWENT CLASS: B04; B05; S03

INVENTOR: BACH F H; BERBERAT P O; ROBSON S C

PATENT ASSIGNEE: (BETH-N) BETH ISRAEL DEACONESS MEDICAL CENT; (BACH-I) BACH F H; (BERB-I) BERBERAT P O; (ROBS-I) ROBSON S C

COUNTRY COUNT: 102

PATENT INFO ABBR.:

PAT	ENT NO	KIN	DATE	WEEK	LA	PG	MAIN	IPC
WO	2003088748	A1	20031030	(200382)*	EN	56[27]		
ΑU	2003226366	A1	20031103	(200438)	EN			
EP	1499186	A1	20050126	(200508)	EN			
JP	2005522521	W	20050728	(200549)	JA	59		
US	20060003922	A1	20060105	(200603)	EN			

APPLICATION DETAILS:

PA:	TENT NO	KIND		APE	PLICATION	DATE
AU EP JP EP JP US	2003088748 2003226366 1499186 A1 2005522521 1499186 A1 2005522521 2006000392 2006000392 2006000392	A1 W W 2 A1 Pr 2 A1	covisional	AU EP JP WO WO US WO	2003-US11411 2003-226366 2003-746978 2003-585506 2003-US11411 2003-US11411 2002-372762E 2003-US11411 2005-511612	20030415 20030415 20030415 20030415 20030415 20030415 20020415 20030415

FILING DETAILS:

PA	TENT NO	KIN	ID	PATENT NO	
EP	2003226 1499186 2005522	A1	Based on Based on Based on	WO 2003088748 WO 2003088748 WO 2003088748	A
PRIORITY	APPLN.		2002-372762P 2005-511612	20020415 20050805	

- L7 ANSWER 2 OF 4 USPATFULL on STN
- TI Spinner preparation machine and cavity resonator
- AB The present invention relates to the treatment of disorders using heme oxygenase-1 and heme degradation products.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.
ACCESSION NUMBER: 2006:4464 USPATFULL

TITLE: Spinner preparation machine and cavity resonator INVENTOR(S):

Bach, Fritz H., Manchester-by-the-sea, MA, UNITED STATES

Berberat, Pascal O., Heidelberg, GERMANY, FEDERAL

REPUBLIC OF

Robson, Simon C., Weston, MA, UNITED STATES

NUMBER KIND DATE -----US 20060003922 A1 20060105 US 2003-511612 A1 20030415 (10) PATENT INFORMATION:

APPLICATION INFO.: WO 2003-US11411 20030415

20050805 PCT 371 date

NUMBER DATE -----

PRIORITY INFORMATION: US 2002-372762P 20020415 (60)

DOCUMENT TYPE: Utility FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, P.O. BOX 1022, MINNEAPOLIS, MN,

55440-1022, US 58

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 27 Drawing Page(s)

LINE COUNT: 3083 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 3 OF 4 USPATFULL on STN

TI Pharmaceutical use of nitric oxide, heme oxygenase-1 and products of heme degradation

AR The present invention relates to the treatment of disorders using nitric oxide (NO), heme oxygenase-1 (HO-1) and heme degradation products such as carbon monoxide (CO), biliverdin, bilirubin and iron.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2004:171542 USPATFULL

TITLE: Pharmaceutical use of nitric oxide, heme oxygenase-1

and products of heme degradation

INVENTOR(S): Bach, Fritz H., Manchester-by-the-sea, MA, UNITED

STATES

Otterbein, Leo E., New Kensington, PA, UNITED STATES

NUMBER KIND DATE PATENT INFORMATION: US 20040131703 A1 20040708 APPLICATION INFO.: US 2003-600182 A1 20030620 (10)

NUMBER DATE PRIORITY INFORMATION: US 2002-390457P 20020621 (60)

DOCUMENT TYPE: Utility

FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, 225 FRANKLIN ST, BOSTON, MA,

02110 23 NUMBER OF CLAIMS:

EXEMPLARY CLAIM: NUMBER OF DRAWINGS: 8 Drawing Page(s)

LINE COUNT: 2300

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 4 OF 4 USPATFULL on STN

TI Methods of treating ileus

AB The present invention relates to a method of treating ileus in a patient, which includes administering a pharmaceutical composition that includes carbon monoxide to the patient.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2003:311902 USPATFULL

TITLE: Methods of treating ileus

INVENTOR(S): Otterbein, Leo E., New Kensington, PA, UNITED STATES
Choi, Augustine M. K., Pittsburgh, PA, UNITED STATES
Moore, Beverley A., Pittsburgh, PA, UNITED STATES
Bauer, Anthony J., Pittsburgh, PA, UNITED STATES

NUMBER DATE

2256

PRIORITY INFORMATION: US 2002-372652P 20020415 (60)
DOCUMENT TYPE: Utility

FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, 225 FRANKLIN ST, BOSTON, MA, 02110

NUMBER OF CLAIMS: 24

EXEMPLARY CLAIM: 1
NUMBER OF DRAWINGS: 16 Drawing Page(s)

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

=> d his

(FILE 'HOME' ENTERED AT 18:15:40 ON 06 JUL 2008)

FILE 'MEDLINE, BIOSIS, WPIDS, USPATFULL, HCAPLUS' ENTERED AT 18:15:59 ON 06 JUL 2008

L1 3209 S (INFLAMMATION AND FERRITIN)

L2 72 S L1 AND BILIVERDIN L3 31 S L2 AND DOSAGE

L4 19 S L3 AND (DEXTRAN)

L5 12 S L3 AND (DESTEROXAMINE)

L6 12 S L5 AND L4 L7 4 S L6 AND (COLITIS)

=> s 16 and (atherosclerosis)

L8 6 L6 AND (ATHEROSCLEROSIS)

=> d 18 ti abs ibib tot

L8 ANSWER 1 OF 6 WPIDS COPYRIGHT 2008 THOMSON REUTERS on STN

TI Treating inflammation involves determining the level of heme oxygenase-l activity in response to a stimulus; and administering anti-inflammatory agent, and composition having carbon monoxide, bilirubin, heme oxygenase-l, and/or appoferritin

AN 2008-C63174 [19] WPIDS

AB WO 2008008513 A2 UPAB: 20080318

NOVELTY - Treating inflammation involves determining the patient's level of heme oxygenaes—($\mathrm{HO}-1$) activity, expression, or induction in response to a stimulus, or determining an allele of a

polymorphism in HO-1 promoter; and administering first pharmaceutical composition comprising an anti-inflammatory agent; and a second pharmaceutical composition comprising carbon monoxide, HO-1, bilirubin, iron dextran, apoferritin (if HO-1 activity, expression or induction in response to stimulus is determined to be reduced compared to reference standard, or if the HO-1 promoter comprises specified allele). DETAILED DESCRIPTION - Treating (M1) inflammation

involves determining the patient's level of heme oxygenase-1 (HO-1) activity, expression, or induction in response to a stimulus, or determining an allele of a polymorphism in HO-1 promoter; and administering first pharmaceutical composition comprising an anti-inflammatory agent; and a second pharmaceutical composition comprising carbon monoxide (CO), a CO-releasing compound, HO-1, bilirubin, biliverdin, ferritin, iron, desferoxamine, salicylaldehyde isonicotinoyl hydrazone, iron dextran, and/or apoferritin (if HO-1 activity, expression or induction in response to a stimulus is determined to be reduced compared to a reference standard, or if the HO-1 promoter comprises a specified allele). An INDEPENDENT CLAIM is included for potentiating (M2) the response of a patient to a pharmaceutical agent, involving identifying a first pharmaceutical agent that is potentiated by a second treatment which induces HO-1 or apoferritin or increases the level of expression of HO-1 or apoferritin in the patient by administering a second pharmaceutical composition comprising HO-1, CO, CO-releasing compound, bilirubin, biliverdin , ferritin, iron, desferoxamine, salicylaldehyde isonicotinoyl hydrazone, iron dextran, and/or apoferritin; administering the first pharmaceutical agent; and allowing the second treatment. PHARMACEUTICALS - Preferred Components: The anti-inflammatory agent is selected from statins, adenosine, cyclooxygenase inhibitors, probucol, steroids, or prostaglandins. In method (M2) the first pharmaceutical agent is immunosuppresant.

ACTIVITY - Antiinflammatory; Antiasthmatic; Respiratory-Gen.; Hypotensive; Cardiovascular-Gen.; Vasotropic; Cerebroprotective; Antiarteriosclerotic; Cardiant; Nephrotropic; Uropathic; Hepatotropic; Virucide: Antiangiogenic: Gastrointestinal-Gen.: Antiarthritic: Antirheumatic; Neuroprotective; Dermatological; Immunosuppressive; Cytostatic; Vulnerary; Nootropic; Antiparkinsonian; Hemostatic; Antibacterial; Analgesic; Gynecological; Endocrine-Gen.; Anti-HIV; Antiallergic. The efficacy of heme oxygenase-1 was tested for anti-inflammatory effect. Mouse macrophage cell lines were stably transfected with either heme oxygenase-1 (HO-I) or empty plasmid (NEO). The resulting cell lines were treated with adenosine (100 muM) or 5'-(N-ethylcarboxamido) adenosine (NECA) (10 muM) 30 minutes prior to stimulation with lipopolysaccharide (LPS) (1 ng/ml). After 4 hours, the supernatant of each group was analyzed for tumor necrosis factor (TNF)-I. Overexpression of HO-1 gave approximately1.5 to 2-fold greater inhibition of TNF-I secretion compared to the vector control. Thus overexpression of HO-1 augmented the effect of the anti-inflammatory agents adenosine and NECA on TNF-I, produced by lipopolysaccharide (LPS) activated macrophages. MECHANISM OF ACTION - Heme oxygenase 1 stimulator; Apoferritin

stimulator. No biological data given.

USE - For treating inflammation (particularly associated with asthma, adult respiratory distress syndrome, interstitial pulmonary fibrosis, pulmonary emboli, chronic obstructive pulmonary disease, primary pulmonary hypertension, chronic pulmonary emphysema, congestive heart failure, peripheral vascular disease, stroke, atherosclerosis, ischemia-reperfusion injury, heart attack, glomerulonephritis, conditions involving inflammation of the kidney, infection of the gentiourinary tract, viral hepatitis, toxic hepatitis, cirrhosis, ileus, necrotizing enterocolitis, inflammatory bowel disease, rheumatoid arthritis, multiple sclerosis, systemic lupus erythematosus, cancer,

wounds, Alzheimer's disease, Parkinson's disease, graft versus host disease, and hemorrhagic, septic, or anaphylactic shock); inflammation of the heart, respiratory tract, liver, spleen, brain, joint, skin, gastrointestinal tract and/or kidney (claimed), pain, reproductive disorders, e.g. impotence, premature uterine contractions, premature deliveries and menstrual cramps, amoebic dysentery, pneumonia (bacterial or viral), inflammatory states due to immunodeficiency e.g. due to infection with HIV; hypersensitivities; and for treating unwanted

ADVANTAGE - The second composition induces HO-1 or increasing the level of expression of HO-1; induces apoferritin or increases the level of expression of apoferritin in the patient, and effectively treat

inflammation.

angiogenesis.

2008-C63174 [19] ACCESSION NUMBER: WPIDS

DOC. NO. CPI: C2008-079637 [19]

TITLE: Treating inflammation involves determining the level of heme oxygenase-1 activity in response to a stimulus; and administering anti-inflammatory agent, and composition having carbon monoxide, bilirubin, heme

oxygenase-1, and/or apoferritin

DERWENT CLASS: B05

BACH F H; HASCHEMI A; OTTERBEIN L E INVENTOR:

(BETH-N) BETH ISRAEL DEACONESS MEDICAL CENT PATENT ASSIGNEE:

COUNTRY COUNT:

PATENT INFO ABBR.:

PATENT NO KIND DATE WEEK LA PG MAIN IPC

WO 2008008513 A2 20080117 (200819)* EN 49[4]

WO 2008008513 A3 20080306 (200819) EN

APPLICATION DETAILS:

PATENT NO KIND APPLICATION DATE WO 2008008513 A2 WO 2007-US16032 20070712

ANSWER 2 OF 6 WPIDS COPYRIGHT 2008 THOMSON REUTERS on STN

TI Use of heme oxygenase-1 and heme degradation products for e.g. reducing inflammation, organ transplantation and treating e.g. cellular proliferative disorder

AN 2003-903222 [82] WPIDS

AB WO 2003088748 A1 UPAB: 20060121

PRIORITY APPLN. INFO: US 2006-830480P

NOVELTY - Reducing (M1) inflammation involves:

(1) administration of at least one treatment selected from inducing ferritin;

20060713

- (2) expressing ferritin; and
- (3) administering a pharmaceutical composition (C1) comprising heme oxygenase-1 (HO-1), bilirubin, biliverdin, ferritin,

iron, desferoxamine, salicylaldehyde isonicotinovl hydrazone,

iron dextran, or apoferritin.

DETAILED DESCRIPTION - INDEPENDENT CLAIMS are also included for:

(1) A method (M2) of transplanting an organ by three different ways (M2a, M2b, M2c). (M2a) involves (ia) administration of at least one of the treatments to a donor to enhance survival or function of the organ after the transplantation, (ib) obtaining an organ from the donor, and (ic) transplanting the organ into a recipient. (M2b) involves (iia) administering to an organ of a donor ex vivo at least one of the

treatments, and (iib) transplanting the organ into a recipient. (M2c) involves (iiia) transplanting an organ from a donor into a recipient, and (iiib) administering at least one of the treatments to the recipient; and (2) A method (M3) of performing angioplasty and vascular surgery

(2) A method (M3) of performing angioplasty and vascular surgery involving performing angioplasty and vascular surgery, respectively, and administering at least one of the treatments.

ACTIVITY - Anti-inflammatory; Antiulcer; Antiasthmatic; Tranquiliager; Respiratory-Gen.; Thrombolytic; Hypotensive; Cardiovascular Gen.; Cerebroprotective; Antiarteriosclerotic; Vasotropic; Cardiant; Nephrotropic; Hepatotropic; Virucide; Gastrointestinal Gen.; Antirheumatic; Antiatrhitic; Vulnerary; Neuroprotective; Nootropic; Antiparkinsonian; Immunosuppressive; Antibacterial; Uropathic; Cytostatic; Gynecological.

The anti-inflammatory efficacy of biliverdin was evaluated in an animal model of endotoxic shock. Administration of endotoxin in male Sprague-Dawley rats resulted in lung inflammation, neutrophil accumulation, and increased levels of tumor necrosis factor-alpha (ThF-alpha) in the serum. The rats were administered with biliverdin (50 micromol/kg) 17 hours prior to, and 8 hours after endotoxin administration. Serum level of ThF-alpha was measured by ELISA kits, and total cell counts was determined by differential analysis. The results showed that biliverdin reduced levels of ThF-alpha; levels of neutrophils and protein accumulation in the airspace; and also increased the levels of anti-inflammatory cytokine IL-10.

MECHANISM OF ACTION - None given.

USE - The treatment is useful for reducing and treating inflammation of the heart, lung, liver, spleen, brain skin and kidney; inflammatory condition (e.g. amoebic dysentery, bacillary dysentery, schistosomiasis, campylobacter enterocolitis, yersinia enterocolitis, enterobius vermicularis, radiation enterocolitis, ischaemic colitis, eosinophilic gastroenteritis, ulcerative colitis, indeterminate colitis, and Crohn's disease) localized in the gastrointestinal tract); asthma; adult respiratory distress syndrome; interstitial pulmonary fibrosis; pulmonary emboli; chronic obstructive pulmonary disease; primary pulmonary hypertension; chronic pulmonary emphysema; congestive heart failure; peripheral vascular disease; stroke; atherosclerosis; ischemia-reperfusion injury; heart attacks; glomerulonephritis; nephrotic disorders; infection of the genitourinary tract; viral and toxic hepatitis; cirrhosis; ileus; necrotizing enterocolitis; specific and non-specific inflammatory bowel disease; rheumatoid arthritis, deficient wound healing, Alzheimer's disease, Parkinson's disease, graft versus host disease, and hemorrhagic, septic, or anaphylactic shock; cellular proliferative and differentiative disorder. For reducing the effects of ischemia. For transplanting organs and performing angioplasty or vascular surgery (all claimed). Also for treating other autoimmune diseases and reproductive disorders.

ADVANTAGE - The heme-oxygenase-1 and the heme degradation products attenuate inflammation and suppress the damage associated with ischemia.

ACCESSION NUMBER: 2003-903222 [82] WPIDS

DOC. NO. CPI: C2003-256695 [82] DOC. NO. NON-CPI: N2003-721263 [82]

TITLE: Use of heme oxygenase-1 and heme degradation products for e.g. reducing inflammation, organ

transplantation and treating e.g. cellular proliferative disorder

DERWENT CLASS: B04; B05; S03

INVENTOR: BACH F H; BERBERAT P O; ROBSON S C

PATENT ASSIGNEE: (BETH-N) BETH ISRAEL DEACONESS MEDICAL CENT, (BACH-I) BACH F H; (BERB-I) BERBERAT P O; (ROBS-I) ROBSON S C COUNTRY COUNT: 102

PATENT INFO ABBR.:

PAT	TENT NO	KINI	DATE	WEEK	LA	PG	MAIN	IPC
WO	2003088748	A1	20031030	(200382)*	EN	56 [27]		
AU	2003226366		20031103		EN			
	1499186		20050126		EN			
	2005522521		20050728		JA	59		
US	20060003922	A1	20060105	(200603)	EN			

APPLICATION DETAILS:

APPLICATION DATE					
WO 2003-US11411 20030415					
AU 2003-226366 20030415					
EP 2003-746978 20030415					
JP 2003-585506 20030415					
WO 2003-US11411 20030415					
WO 2003-US11411 20030415					
US 2002-372762P 20020415					
WO 2003-US11411 20030415					
US 2005-511612 20050805					

FILING DETAILS:

PA'	TENT NO	KIND	P	ATENT NO
EP	2003226366 A 1499186 A1 2005522521 V	Based	on W	O 2003088748 A O 2003088748 A O 2003088748 A
PRIORITY	APPLN. INFO:	US 2002-372 US 2005-511		020415 050805

- L8 ANSWER 3 OF 6 USPATFULL on STN
- ΤI Spinner preparation machine and cavity resonator
- AB The present invention relates to the treatment of disorders using heme oxygenase-1 and heme degradation products.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2006:4464 USPATFULL TITLE: Spinner preparation machine and cavity resonator

INVENTOR(S): Bach, Fritz H., Manchester-by-the-sea, MA, UNITED STATES

Berberat, Pascal O., Heidelberg, GERMANY, FEDERAL REPUBLIC OF

Robson, Simon C., Weston, MA, UNITED STATES

NUMBER KIND DATE PATENT INFORMATION:

US 20060003922 A1 20060105 US 2003-511612 A1 20030415 20030415 (10) APPLICATION INFO.: WO 2003-US11411 20030415

20050805 PCT 371 date

NUMBER DATE PRIORITY INFORMATION: US 2002-372762P 20020415 (60) DOCUMENT TYPE: Utility

APPLICATION FILE SEGMENT:

LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, P.O. BOX 1022, MINNEAPOLIS, MN,

55440-1022, US

NUMBER OF CLAIMS: 58 EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 27 Drawing Page(s)

LINE COUNT: 3083

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L8 ANSWER 4 OF 6 USPATFULL on STN

TΙ Methods of treating angiogenesis, tumor growth, and metastasis

AB The present invention relates to a method of treating cancer or unwanted angiogenesis in a patient, which includes administering a pharmaceutical composition that includes carbon monoxide to the patient.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2004:326952 USPATFULL

Methods of treating angiogenesis, tumor growth, and TITLE:

metastasis

Otterbein, Leo E., New Kensington, PA, UNITED STATES INVENTOR(S):

Choi, Augustine M. K., Pittsburgh, PA, UNITED STATES

NUMBER KIND DATE PATENT INFORMATION: US 20040258772 A1 20041223 US 2003-455564 A1 20030605 (10)

APPLICATION INFO.:

AB

NUMBER DATE

_____ PRIORITY INFORMATION: US 2002-386561P 20020605 (60)

DOCUMENT TYPE: Utility

FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, 225 FRANKLIN ST, BOSTON, MA, 02110

NUMBER OF CLAIMS: 71 EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 10 Drawing Page(s)

LINE COUNT: 1303

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 5 OF 6 USPATFULL on STN

ΤI Pharmaceutical use of nitric oxide, heme oxygenase-1 and products of heme degradation

The present invention relates to the treatment of disorders using nitric oxide (NO), heme oxygenase-1 (HO-1) and heme degradation products such

as carbon monoxide (CO), biliverdin, bilirubin and iron.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2004:171542 USPATFULL

TITLE: Pharmaceutical use of nitric oxide, heme oxygenase-1

and products of heme degradation

INVENTOR(S): Bach, Fritz H., Manchester-by-the-sea, MA, UNITED

STATES

Otterbein, Leo E., New Kensington, PA, UNITED STATES

NUMBER KIND DATE PATENT INFORMATION: US 20040131703 A1 20040708 APPLICATION INFO.: US 2003-600182 A1 20030620 (10) NUMBER DATE

PRIORITY INFORMATION: US 2002-390457P 20020621 (60)

02110

DOCUMENT TYPE: Utility

FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, 225 FRANKLIN ST, BOSTON, MA.

NUMBER OF CLAIMS: 23

EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 8 Drawing Page(s) LINE COUNT: 2300

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 6 OF 6 USPATFULL on STN

ΤI Methods of treating vascular disease

AB The present invention relates to a method of treating patients suffering from, or at risk for, intimal hyperplasia and/or arteriosclerosis. The treatment includes administering a pharmaceutical composition that includes carbon monoxide to the patient.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER: 2003:311901 USPATFULL

TITLE: Methods of treating vascular disease

Otterbein, Leo E., New Kensington, PA, UNITED STATES INVENTOR(S): Choi, Augustine M. K., Pittsburgh, PA, UNITED STATES Bach, Fritz H., Mancester-by-the-sea, MA, UNITED STATES

Zuckerbraun, Brian, Pittsburgh, PA, UNITED STATES

NUMBER KIND DATE PATENT INFORMATION:

US 20030219496 A1 20031127 US 7364757 B2 20080429 US 2003-367277 A1 20030213 (10) APPLICATION INFO.:

> NUMBER DATE -----

PRIORITY INFORMATION: US 2002-356718P 20020213 (60)

DOCUMENT TYPE: Utility FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, 225 FRANKLIN ST, BOSTON, MA,

02110

NUMBER OF CLAIMS: 49 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 18 Drawing Page(s)

LINE COUNT: 1841

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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(FILE 'HOME' ENTERED AT 18:15:40 ON 06 JUL 2008)

FILE 'MEDLINE, BIOSIS, WPIDS, USPATFULL, HCAPLUS' ENTERED AT 18:15:59 ON 06 JUL 2008

3209 S (INFLAMMATION AND FERRITIN) L1 L2 72 S L1 AND BILIVERDIN

L3 31 S L2 AND DOSAGE

19 S L3 AND (DEXTRAN) T. 4

L5 12 S L3 AND (DESFEROXAMINE)

1,6 12 S L5 AND L4 T.7 4 S L6 AND (COLITIS) =>